In the Claims:

Please amend the claims as shown:

Claims 1-7. (Canceled)

8. (Previously Amended) An isolated antibody to an isolated peptide of the general formula:

G-XI-X2-R (SEQ ID NO: 1)

and wherein G is glutamate or glutamine;

X1 is a bond or an amino acid selected from the group consisting of isoleucine, valine, methionine, alanine, and phenylalanine;

X2 is an amino acid selected from the group consisting of aspartate, glutamate, and asparagine; and

R is a tripeptide wherein each of the amino acids of said tripeptide are selected from the group consisting of proline, isoleucine, aspartate, alanine, glycine, serine, valine, lysine, glutamate, glutamine, threonine, leucine, methionine, phenylalanine, arginine, and asparagine;

and wherein the antibody binds to a 6C5 antigen.

9. (Original) The antibody according to claim 8, wherein said antibody is a monoclonal antibody.

10. (Original) The antibody according to claim 8, wherein said antibody is 6C5.

11. (Original) The antibody according to claim 8, wherein said antibody is produced by the cell line F6-6C5-H4 (ATCC No. PTA-1358).

12. (Original) The antibody according to claim 8, wherein said antibody is a polyclonal antibody.

13. (Original) The cell line F6-6C5-H4 (ATCC No. PTA-1358).

14. (Original) A method of treating a yeast infection in a patient in need of such treatment comprising administering to said patient a composition comprising an active agent, wherein said active agent is an antibody according to claim 8.

15. (Original) A method of detecting a hydrophobic binding domain in a sample containing multiple components, comprising the steps of:

providing an antibody according to claim 8, wherein said antibody is labeled with a detectable marker;

contacting the sample with an antibody according to claim 8; and

isolating any resulting complexes formed between the sample components and the labeled antibodies.

16. (Original) The method of claim 15, wherein said detection is performed in vivo.

17. (Original) The method of claim 15, wherein said detection is performed in vitro.

18. (Original) A method of isolating a hydrophobic binding domain comprising the steps of:

providing an antibody according to claim 8, wherein said antibody is labeled with a detectable marker, and wherein said antibody is bound to a solid support;

contacting a sample containing multiple components with said antibody; and

washing the solid support to remove unbound material.

Claims 19-48. (Canceled).

49. (Currently Amended) The antibody of claim 8, wherein said peptide comprises Glutamate-Isoleucine-Aspartate-Proline-Isoleucine-Aspartate **(SEQ ID NO: 41)**.

50. (Currently Amended) The antibody of claim 9 8, wherein the peptide consists of Glutamate-Isoleucine-Aspartate-Proline-Isoleucine-Aspartate **(SEQ ID NO: 41)**.

51. (Original) The antibody of claim 49, wherein the antibody is a monoclonal antibody.

52. (Original) The antibody of claim 50, wherein the antibody is a monoclonal antibody.

53. (Original) The antibody of claim 49, wherein the antibody is a polyclonal antibody.

54. (Original) The antibody of claim 50, wherein the antibody is a polyclonal antibody.